

63. The method of claim 53, wherein the ligating comprises ligating to an overhang or a blunt end.

64. The method of claim 53, further comprising detecting mutations in one or more of the error-corrected sequences as compared to a reference sequence.

65. The method of claim 53, wherein sequencing the cypher-target amplification products comprises converting data from a sequencing instrument into quality scores and then into sequencing reads.

66. The method of claim 53, further comprising purifying a plurality of cypher-target nucleic acid complexes prior to sequencing, wherein the purified cypher-target nucleic acid complexes comprise nucleic acid molecules from specific genomic regions.

67. The method of claim 53, wherein the plurality of circulating DNA molecules comprise a mutation present at a frequency of 2.1×10^{-6} or lower.

68. The method of claim 53, wherein generating the error corrected sequences results in a measureable sequencing error rate from about 10^{-6} to about 10^{-8} .

69. The method of claim 53, wherein the circulating DNA molecules comprise plasma DNA biomarkers.

70. The method of claim 53, wherein each identifier tag of the plurality of distinct identifier tag sequences is a random or partially random sequence of about 5 nucleotides in length.

71. The method of claim 53, wherein each identifier tag of the plurality of distinct identifier tag sequences is a random or partially random sequence of 5 or 6 nucleotides in length.

72. The method of claim 53, wherein the cypher polynucleotides comprising the identifier tags are contained within a pool of cypher polynucleotides comprising known sequences.

73. The method of claim 53, wherein the ligating comprises ligating identifier tags to both ends of the circulating DNA molecules, and further wherein the identifier tags at both ends together form a unique pair of identifiers that differ between each of the other pairs of identifiers ligated to the circulating DNA molecules.

74. The method of claim 53, wherein grouping sequencing reads is based on (i) the identifier tag sequence and (ii) sequence information from an end of the circulating DNA molecule.

75. The method of claim 61, further comprising purifying a plurality of cypher-target nucleic acid complexes prior to sequencing, wherein the purified cypher-target nucleic acid complexes comprise nucleic acid molecules from specific genomic regions.

76. The method of claim 61, further comprising identifying one or more single nucleotide mutations.

77. The method of claim 61, wherein the circulating DNA molecules comprise blood biomarkers.

78. The method of claim 61, wherein the circulating DNA molecules comprise DNA molecules derived from cancer cells.

79. The method of claim 61, wherein each identifier tag of the plurality of distinct identifier tag sequences is a random or partially random sequence of about 5 nucleotides in length.

80. The method of claim 61, wherein each identifier tag of the plurality of distinct identifier tag sequences is a random or partially random sequence of 5 or 6 nucleotides in length.

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